

TEST CHANGE

Paraneoplastic Pemphigus (Paraneoplastic Autoimmune Multiorgan Syndrome) Antibody Screening Antibodies by IIFPanel

0092107, PARA PEMPH

Specimen Requirements:

Patient Preparation:

Collect: Plain red or serum separator tube (SST).

Specimen Preparation: Transfer 2 mL serum to an ARUP <u>standard transport</u>

tube. Standard Transport Tube. (Min: 0.5 mL)

Effective Date: August 21, 2023

Transport Temperature: Refrigerated.

Unacceptable Conditions: Hemolyzed or lipemic specimens. Plasma.

Remarks:

Stability: Ambient: 1 week; Refrigerated: 2 weeks; Frozen: Indefinitely

Methodology: Semi-Quantitative Indirect Immunofluorescence (IIF)

Performed: Varies

Reported: 3-7 days

Note: The methodology is indirect immunofluorescence (IIF) of

patient serum on substrates from rodents including rat bladder, mouse bladder, mouse heart, and mouse liver to detect characteristic antibody reactivity: simple columnar epithelial cell surface and basement membrane zone in bladders, intercalated discs in heart, and portal tracts in liver. Monkey esophagus substrate is included if other concurrent IIF testing

does not. For specimens less than 0.5 mL, call the

Immunodermatology Laboratory at (866₋)-266-5699. This test should be distinguished from antibody testing of cerebral spinal fluid (CSF) for paraneoplastic neurologic syndromes;

3004510, 3004512, 3004517 are different tests.

CPT Codes: 88346; 88350 x4

New York DOH Approval Status: This test is New York DOH approved.

Interpretive Data:

Refer to report

Reference Interval:



Effective Date: August 21, 2023

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HOTLINE NOTE: There is a component change associated with this test. One or more components have been added or removed. Refer to the Hotline Test Mix for interface build information.